IN THE CLAIMS

1. (original) Uracilo A uracil having general formula (I):

(I)

wherein:

3 X₁ represents a hydrogen atom or a halogen atom;

- 4 X2 represents a halogen atom;
- X4 represents a C1-C3 haloalkyl group;
- R represents a hydrogen atom, a C₁-C₃ alkyl group or a C₁-C₃ haloalkyl group;
- G represents an oxygen atom or a sulphur atom;
- X₃ represents a Q(CR₁R₂)_nZ- group, a Q₂-Z- group, a Q₂-group, a Y(OC)-CR₆=CR₅-CR₃R₄Z- group;
- Z represents an oxygen atom or a sulphur atom;
- R₁, R₂, R₃ and R₄, the same or different, represent a hydrogen atom or, a C₁-C₄ alkyl group; or a C₂-C₄ haloalkyl group;
- R₅ represents an OR₇ group;
- Re represents a hydrogen atom or a C1-C4 alkyl group;
- R₇ represents a C₁-C₄ alkyl group or a C₁-C₄ haloalkyl group;
- Y represents a C4-C6 alkoxy or haloalkoxy group; an OR4
 group, a SRa group, a NRaRu eroup;

turn, possibly substituted with one or more halogen atoms selected from chlorine, fluorine, bromine or todine, or substituted with one or more groups selected from C₄-C₄ alkyl, or C₄-C₄ haloslkyl, C₄-C₄ alkony or C₄-C₄ haloslkony.

Ale-and Rill, the same or different, represent a hydrogen atom, or a C₂-C₆ alkyl group, a C₂-C₆ halcalkyl group, or an aryl group, each cycloalkyl group, or an aryl group, each groups, in turn, possibly substituted with one or more halogen atoms selected from chlorine, fluorine, bromine or iodine, or substituted with one ox more groups selected from a C₄-C₄ alkyl, or C₄-C₅ halcalkyl, C₄-C₄ alkowy or C₄-C₄ halcalkony, or, jointly represent a C₄-C₄ alkylene chain possibly substituted with C₄-C₄ alkyl groups and possibly interrupted by charges atoms or by a NR₃, group, wherein.

- Ris represents a hydrogen atom, a Cs-Cs-olkyl group or Cs-Cs-olkenyl group or a Cs-Cs-olkenyl group or a Cs-Cs-olkenyl group or a Cs-Cs-olkylcarbonyl group or Cs-Cs-olealkylcarbonyl group
- n represents 1, 2 or 3;
- Q represents a heterocyclic group selected from pyrrol-

2-yl, pyrrol-3-yl, imidasol-2-yl, imidasol-4-yl, imidazol-5-yl, pyrazol-3-yl, pyrazol 4-yl, pyrazol 6yl, 1,2,4-triazol-3 yl, 1,2,4-triazol-5-yl, 1,2,4triazol-3-onyl, -1,2,3-triazolyl, tetrazolyl, oxazolyl, isomazol-5-yl, thiazol-2-yl, thiazol-5-yl, isothiazolyl, 1,3,4-oxadiazolyl, 1,3,4-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,4-oxadiazolyl, 1,2,4-oxadiazol 5-on-3-yl, benzoxazol-2-yl, benzothiazol-2-yl, pyrasinyl, pyridazinyl, 1,2,4 triozinyl, 1,3,4 thiadiazol-2-on-5-yl, 1,4,2-dioxazol-5-on-3-yl, 1,4,2oxathiasol-5-on-3-yl, 1,3,4-oxadiazin-5-on-2-yl, 1,4,2dioxazin-3-yl, --1,2,4-oxadiazin-5-on 3-yl, 4,5,6,7tetrahydre-1,3-benzothiazol-2-yl, 5,6-dihydro-4Heyelopenta[d][1,3]thiazole, said groups, in turn, possibly being optionally substituted with a halogen atom[[s]] selected from chlorine, fluorine, bromine or iodine, or substituted with a group[[s]] selected from C_1-C_6 alkyl or C_1-C_6 haloalkyl, C_2-C_6 alkenyl or C_2-C_6 haloalkenyl, Ga-Ga alkenylony or Ga-Ga haloalkenylony. C2-C6 alkynyl or C2-C6 haloalkynyl, G2-C5 alkynyloxy or C. C. haloalkynyloxy, C. C. alkoxy or C. C. haloalkoxy. Ga-Ga alkowyalkyl or Ga-Ga haloalkowyalkyl, Ga-Ga alkoxyalkoxy, C2-C6 holoalkoxyalkoxy, C2-C6 halealkeжyhalealkeжy, С₃-С₆ alkeжyalkeжyalkyl, С₃-С₉

alkoxyalkoxyalkoxy, C1-C4 alkylthio or C1-C4 haloalkylthio, -Ca-Ca alkylthioalkyl, -Ca-Ca-alkyloulfinie or C1-C4 haloalkyloulfinic, C1-C4 alkyloulfonic or C1-C4 haloalkylsulfonic, C. G. alkoxycarbonyl or C. G. haloalkoxycarbonyl, C₄-C₁ alkenyloxycarbonyl or C₄-C₄ alkynyloxyearbonyl, C,-C,-C, alkoxyearbonylalkyl or C,-C, haloulkoxycarbonylalkyl, C4-C9_alkenyloxycarbonylalkyl -C₄-C₃ alkynyloxyearbonylalkyl, C₂-C₃ alkowycarbonylalkowy, C4-G2 alkenylowycarbonylalkowy or ___alkynylemysarbonylalkemy, aminocarbonylalkoxy possibly substituted with C+-C4 alkyl groups or with a C2-C5 alkylene group, CN, CHO, NO27 NH27 OH, C1-C2 cyanoalkyl, C1-C3 cyanoalkylony, C2-C6 formylalkyl, C2-C6 alkylearbonyl, C2-C6 haloolkylcarbonyl, Ca-Ca alkylcarbonylalkyl, Ca-Ca alkoxvimino, Ca-C6 haloalkonyimino, Ca-C6 alkoxyiminoalkyl, C. G. haloalkoxyiminoalkyl, C.-C. alkoxyiminehalealkyl, aminecorbonyl, C2-C4 eminocarbonylalkyl, aminosulfonyl or C2-C5 aminosulfonylalkyl, these last four groups possibly oubstituted-with one or two C, C, alkyl-groups or with a G2-G5 alkylene group; G1-G6 alkyloulfonylamine, G2-G7 alkylcarbonylamino or C2-C2 alkoxycarbonylamino, these last three groups possibly substituted with G.-C. alkyl

2. represente a heterocyclic group oclected from 1,3,4-thiadiczol 2 yl, 1,3,4-thiadiczol 5 yl, 1,2,4-thiadiczol 5 yl, tetracel 5 yl, 1,3,4-enadiczol 2 yl, 1,3,4-enadiczol 2 yl, 1,3,4-enadiczol 2 yl, 1,3,4-enadiczol 5 yl, exazol 2 yl, enazol 3 yl, isomazol 3 yl, isomazol 5 yl, thiczol 3 yl, thiazol 5 yl, thiazol 5 yl, thiazol 5 yl, caid groups, in turn, possibly substituted with halogen atoms solected from chlorine, fluorine, bromine ox iodine, or substituted with groups solected from C₂-C₆ alkyl or C₂-C₆ haloalkyl, C₃-C₆ alkenyl or C₂-C₆ haloalkynyl or C₂-C₆ haloalkynyl, C₃-C₆ alkenyl or C₄-C₆ haloalkynylony, C₅-C₆ alkonyl or C₅-C₆ haloalkynylony, C₅-C₆ alkonyl or C₅-C₆ haloalkynyl, C₅-C₆ alkonyl or C₅-C₆ haloalkynyl, C₅-C₆ alkonylyl, C₅-C₆ alkonylony, C₅-C₆ alkonylyl, C₅-C₆ alkonylonyl, C₅-C₆ alkonylonyl, C₅-C₆ alkonylonyl, C₅-C₆ alkonylonyl, C₅-C₆ alkyloulfinic or C₆-C₆ haloalkonyl, C₅-C₆ alkyloulfinic or

C.-C. haloalkyloulfinic, C.-C. alkyloulfonic-or C.-C. haloalkyloulfonic, -C2 -C4 alkoxycarbonyl or C2-C4 haloalkoxycarbonyl, C2-C2 alkoxycarbonylalkyl or C2-C2 haloalkoxycarbonylalkyl, Ga-Ca alkoxycarbonylalkoxy, Ca-Ga aminocarbonylalkoxy-possibly-substituted with C1-G4 alkyl groups or with a G2-G5 alkylene; CN, CHO, NO2, NH2, -C1-C2 cyanoalkyl, -C1-C2 cyanoalkyloxy, -C2-C4 alkylcarbonyl, C. C. haloalkylcarbonyl, C. C. alkoxyiminoalkyl, Ca-Ca haloalkoxyiminoalkyl, aminocarbonyl, C. C. aminocarbonylalkyl, aminosulfonyl o C2-C6 aminosulfonylalkyl, these last four groups possibly substituted with one or two-C+-C+-alkyl groups or with a Ca-Ca alkylene, Ca-Ca alkyloulfonylamino, Ca-Ca alkylcarbonylamino or C2-C2 alkoxycarbonylamino, these last three groups possibly substituted with G, C, alkyl groups, C. C10 aryl, C4 C12 arylalkyl, C4 C10 arylalkoxy, C+-G12 aryloxyalkyl, Ge-G12 arylalkyloxyalkyl said groups in turn possibly substituted with halogen atoms, C.-C. alkyl-groupe,-C,-C,-holoalkyl-groupe,-C,-C,-alkeny groups, C1-C2 haloalkoxy groups, CN+-C2-C2-cycloalkyl, C. C12 cycloalkylalkyl, C. C10 cycloalkylalkony, tetrahydropyran-2-yl said groups in turn-possibly oubstituted with halogen atomo, C+-C4-alkyl groups, C+-G4_alkoxy groups;

 Q_2 represents a heterocyclic group selected from 1Htetrazol-5-yl or 2H-tetrazol-5-yl, thiazol-2-yl, thiasel-4-yl, thiasel-5-yl, icothiasel-3-yl, isothiazol 4-yl, isothiazol 5-yl, 1,2,3-triazolyl, benzoxazol 2-yl, benzothiazol-2-yl, pyrimidin-2-yl, 1,2,4-triazinyl,-1,3,5-triazinyl,-1,3,4-thiadiazol-2on-5-yl, 1,4,2-dioxazol-5-on-3-yl, -1,4,2-oxathiasol-5on-3-yl, 1,3,4-oxadiazin-5-on-2-yl, 1,4,2-dioxazin-3yl, 1,2,4-omadiazin-5-on-3-yl, 4,5,6,7-totrahydro-1,3benzethiazel-2-yl, -5,6-dihydro-4Heyelopenta(d)(1,3)thiazole, said-groups in turn possibly being optionally substituted with halogen atoms selected from chlorine, fluorine, bromine or iodine, or substituted with a group[[s]] selected from: C_1-C_6 alkyl; [[or]] C_1-C_6 haloalkyl[[,]]; C_2-C_6 alkenyl; [[or]] C2-C6 haloalkenyl[[,]]; G2-C6 alkenylony or C2-C6 haloalkenyloxy, C2-C6 alkynyl; [[or]] C2-C6 haloalkynyl, C2-C4 alkynyloxy-or-C2-C4 haloalkynyloxy, C4-C4 alkoxy or G:-C: haloalkowy, C2-C: alkoxyalkyl; [[or]] C2-C: haloalkoxyalkyl[[,]]; Ca-Ca alkoxyalkoxy, Ca-Ca haloalkoxyalkoxy, C2-Cs haloalkoxyhaloalkoxy, C3-C8 alkoxyalkoxyalkyl, G. C. alkonyalkoxyalkoxy, C. G. $alkylthio or G_1-G_6$ haloalkylthio, G_2 - G_6 alkylthioalkyl, G1-G4-alkyloulfinie-or-C4-C4-haloalkyloulfinie,-C4-C4

alkylsulfonic-or-C₂-C₆ haloalkylsulfonic,-C₂-C₆ alkonycarbonyl or C.-C. halealkonycarbonyl, C. C. alkenyloxyearbonyl or C2-C2 alkynyloxycarbonyl, C2-C2 alkoxycarbonylalkyl or G. G. haloalkoxycarbonylalkyl, G4-G2 alkenylonycarbonylalkyl or C4-C4 alkynylexycarbonylalkyl, G2-C4-slkonycarbonylalkoxy, alkenylonyearbonylalkeny C4-G2 or alkynyloxycarbonylalkoxy C4-C4, C4-C4 aminocarbonylalkoxy possibly substituted with C. C. alkyl or with a Ca-Ca alkylone; CN, GHO, NO2, NH2, OH, C1-C3 Gyanoulkyl, C1-C3 Gyanoulkyloxy, C2-C6 formylalkyl, Ca-Ca alkylearbonyl, Ca-Ca haloalkylearbonyl, Ca-Ca alkylcarbonylalkyl, C2-C4 alkoxyimino, C2-C4 haloalkonyimino, Ca-Calkonyiminoalkyl, Ca-Ca haloalkonyiminoalkyl, alkonyiminohaloalkyl Ca-Caaminocarbonyl, G2-Gc aminocarbonylalkyl, aminosulfonyl er-Ca-Ca-aminosulfonylalkyl, these last four groups possibly substituted with one or two C1-C1 alkyl groups er with a G2-G4-alkylene; G1-C4-alkyleulfonylamine, G2-G1 alkylcarbonylamino o C2-C2-alkonycarbonylamino, these last three-groups possibly substituted with C. C. alkyl groups; C. C. aryl, C. C. arylalkyl, C. arylalkowy, C7-C12 aryloxyalkyl, C8-C12 arylalkyloxyalkyl said groups in turn pessibly being optionally substituted with

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halogen atoms, C1-C4 alkyl groups, C1-C3 haloalkyl
     groups, C1-C4 alkoxy groups, C1-C3 haloalkoxy groups, CN;
     C3-C7 cycloalkyl, C6-C12 cycloalkylalkyl, C6-C10
     cycloalkylalkory, tetrahydropyran-2-yl said groups in
     turn possibly being optionally substituted with halogen
     atoms, C1-C4 alkyl groups, C1-C4 alkoxy groups.
2. (original): The uracils A uracil according to claim 1,
characterized in that they are it is selected from:
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-
3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-(2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-(2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenylthio}-3-methoxybut-2-enoate;
- ethyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
vllphenoxy}-3-ethoxybut-2-enoate;
- methyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenylthio}-3-methoxybut-2-enoate;
- ethyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
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methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-ethoxybut-2-enoate;

    isopropyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-

tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-
1-y1]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
methoxybut-2-enoate;
- methyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy)-3-
methoxybut-2-enoate;
- ethyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
ethoxybut-2-enoate;
- othyl (25)-4-[2,4-dichloro-5-[1,2,3,6-tetrahydro-2,6-
dioxo-4-(trifluoremethyl)pyrimidin-1-yl]phenoxy)-3-
ethoxybut-2-eneate;
-2,2,2-trifluoreethyl (2E)-4-(2-chlore-4-fluore-5-
{1,2,3,6-tetrahydre-2,6-dioxo-4-(trifluoromethyl)pyrimidin-
1-yl]phenoxy)-3-methoxybut-2-enoater
- (2E) -1 (2 chlore 1-fluore-5-[1,2,3,6-tetrahydre-2,6-
dioxo-4 (trifluoromethyl)pyrimidin-1-yl]phenoxy)-3-methoxy-
N, N-dimethylbut-2-enamide;
- S-cthyl-(2E)-4-(2-chloro-4-fluoro-5-(1,2,3,6-tetrahydro-
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2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy)-3-
methoxybut-2-enethicater
- isopropyl (2E)-4-(2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
vl]phenoxy}-3-methoxybut-2-enoate;
- 2,2,2-trifluoroethyl (2E)-4-{2-chloro-4-fluoro-5-
[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluorome-
thyl)pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- 2,2,2-trifluoroethyl (2E)-4-(2,4-dichloro-5-[1,2,3,6-
tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)
pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- 5-ethyl-(2E)-4-(2-ehlero-4-fluoro-5-(1,2,3,6-tetrahydro-
3-methyl-2,6-dioxe-4-(trifluoremethyl)-pyrimidin-1-
yl]phonoxy)-3-motherybut-2-enethicater
- S-ethyl-(2E)-4-(2,4-dichlore-5-[1,2,3,6-tetrahydre-3-
mothyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yllphenoxy)-2-methoxybut-2-enethicater
-- (2E)-4-(2-chloro-4-fluoro-5-(1,2,3,6-tetrahydro-3-methyl-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phonoxy)-3-
methoxy-N, N-dimethylbut-2-enamide:
- (2E)-4-(2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-methoxy-
N-N-dimethylbut-2-enamider
-- (2E)-4-{2-chloro-4-fluoro-5-{1,2,3,6-tetrahydro-3-methyl-
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2,6-dioxe-4-(trifluoremethyl)pyrimidin-1-yl]phonylthio)-3-
methoxy N, N-dimethylbut-2-enamide;
- (2E)-4-(2,4-dichlore-5-[1,2,3,6-tetrahydre-3-methyl-2,6-
diexe-4-(trifluoromethyl)pyrimidin-1-yl]phenylthie)-3-
methoxy-N,N-dimethylbut-2-enamider
- 3-{4-ohloro-2-fluoro-5-(tetrazol-5-ylmethoxy)phonyl}-6-
(trifluoromethyl)-2, 4 (1H, 3H)-pyrimidinodione+
- 3-(4-ohloro-2-fluoro-5-[(2-methyl-2H-tetrazol-5-
yl)methonyl)-6-(trifluoromethyl)-2,4(1H,3H)
pyrimidinedione.
-3-[4-chloro-2-fluoro-5-(tetrazol-5-ylmothoxy)phcnyl]-1-
methyl-6 (trifluoromethyl) 2,4(1H,3H)-pyrimidinedione+
-3-[2,4-dichloro-5-(tetrazol-5-ylmethoxy)phonyl]-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyximidinedione;
- 3-(4-chloro-2-fluoro-5-((2-methyl-2H-tetrozol-5-
y1) methony | pheny1 | - 1 - methyl - 6 - (trifluoromethyl) - 2, 4 (1H, 3H) -
pyrimidinedione:
- 3-[4-chloro-2-fluoro-5-[(2-cthyl-2#-tetrazol-5-
yl)methoxy]phenyl]-1 methyl-6-(trifluoremethyl)-2,4(1H,3H)-
pyrimidinediene;
-3-12, 4-dichlore-5-[(2-methyl-2H tetrazel-5-
y1) methoxy | phenyl | -1 -methyl -6 -(trifluoremethyl) -2, 4 (1H, 3H) -
pyrimidinodioner
- 3-12,4-dichloro-5-[(2-ethyl-2H-tetrazol-5-
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y1)methoxylphenyl) 1-methyl 6-(trifluoromethyl)-2,4(1H,3H)
pyrimidinediene:
-3-(4-chlore-2-fluore-5-(1-cthyl-1#-tctrasel-5-
yl)methoxylphonyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione:
-3-(2,1-dichlore-5-[(1-cthyl-1#-tctrare1-5-
yl) methoxylphenyl) -1-methyl-6-(trifluoromethyl) -2,4(1H,3H)-
pyrimidinedione;
- 3-{5-[(5-tert-butyl-1,3,4-oxadiazol-2-y1)methoxy]-4-
chloro-2-fluorophenyl}-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H)-pyrimidinedione;
- methyl [5 ((2-chloro-4-fluoro-5-(1,2,3,6-tetrahydro-3-
methyl-2,6-diono-4-(trifluoromethyl)pyrimidin-1-
yl)phonoxy|methyl)-1H-tetrazol-1-yl]acetate;
--mothyl [5-({2,4-dichlore-5-[1,2,3,6-tetrahydro-3-methyl-
2.6 dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phonoxy;methyl)-1H-tetrazol-1-yl]acetate;
- mothyl [5-((2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phonoxy|methyl) 2H-tetrazel-2-yl]acetate;
- mothyl [5-((2,4-dichlore-5-[1,2,3,6-tetrahydre-3-methyl-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy|methyl)-2H-tetrazol-2-yl]acetate+
- 3-[4-chloro-3-(tetrazol-5-yl)phenyl]-6-(trifluoromethyl)-
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2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-3-(2-methy1-2H-tetrazo1-5-y1)pheny1]-6-
 (trifluoromethy1)-2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-3-(1-methyl-1H-tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-3-(tetrazol-5-yl)phenyl]-1-methyl-6-
 (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(tetrazol-5-yl)phenyl]-6-
- (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[2,4-dichloro-5-(tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(tetrazol-5-yl)phenyl]-1-methyl-6-
(trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[2,4-dichloro-5-(tetrazol-5-yl)phenyl]-1-methyl-6-
(trifluoromethyl) -2,4(1H,3H)-pyrimidinedione:
- 3-[4-chloro-3-(2-methy1-2H-tetrazol-5-y1)pheny1]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione:
- 3-[4-chloro-2-fluoro-5-(2-methyl-2H-tetrazol-5-
yl)phenyl]-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[2,4-dichloro-5-(2-methyl-2H-tetrazol-5-yl)phenyl]-6-
(trifluoromethy1)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(1-methyl-1H-tetrazol-5-
yl)phenyl]-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[2,4-dichloro-5-(1-methyl-1H-tetrazol-5-yl)phenyl]-6-
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(trifluoromethy1) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(2-methyl-2H-tetrazol-5-
yl) phenyl] -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H) -
pyrimidinedione;
- 3-[2,4-dichloro-5-(2-methyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-(4-chloro-3-(2-ethyl-2H-tetrazol-5-yl)phenyl]-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-3-(1-methyl-1H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(1-methyl-1H-tetrazol-5-
yl)phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-[2,4-dichloro-5-(1-methyl-1H-tetrazol-5-vl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-3-(1-ethyl-1H-tetrazol-5-yl)phenyl]-1-methyl-
6-(trifluoromethy1)-2,4(1H,3H)-pyrimidinedione;
- methyl (5-[2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-
dioxe-4-(trifluoromethyl)pyrimidin-1-yl]phenyl} -1#-
totrazol-1-yl)acetate;
-- mothyl-(5-{2-chloro-5-{1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl-phenyl)-2H-
tetrazel 2 yl) acetate;
- methyl-(5-(2-chlore-4-fluore-5-(1,2,3,6-tetrahydre-3-
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methyl-2,6-dioxe-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1H-tetrazel-1-vl)acetater - mothyl (5 (2-chloro 4-fluoro 5-[1,2,3,6-tetrahydro-3methyl-2,6-diono-4-(trifluoromethyl)pyrimidin-1-yl]phenyl)-2H-tetrasel-2-yl-acetate; - methyl (5-(2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxe-1-(trifluoremethyl)pyrimidin-1-yl]phenyl)-1Htetrazol-1-yl)acetate; -- methyl (5-(2,4-dichlero-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-vl]phenyl)-2Htetrazel 2 vllacetate. -3-[4-chloro-3-(4-methoxy-5-methyl-1,3-thiazol-2yl) phenyl-6-(trifluoromethyl)-2, 4(1H, 3H)-pyrimidinedione; - 3-[2,4-dichloro-5-(4-methoxy-5-methyl-1,3-thiazol-2v1) phonyl 6-(trifluoromethyl)-2, 4(1H, 3H)-pyrimidinedione, - 3-[4-chloro-2-fluoro-5-(4-methoxy-5-methyl-1,3-thiazol-2y1)phonyl-6-(trifluoromethyl)-2,4(1H,3H)-pyximidinodioner --3-[4-chloro-3-(4-methoxy-5-methyl-1,3-thiazel-2yl) phonyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)pyrimidinediene: - 3-[4-chloro-3-(4-cthoxy-5-methyl-1,3-thiazo1-2-yl)phenyl-1-mothyl-6-(trifluoromethyl) 2,4(1H,3H)-pyrimidinodione+ - 3-[2, 4-dichloro-5-(4-methoxy-5-methyl-1,3-thiazel-2v1) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-

```
pyrimidinediene;
- 3-[2,4-dichlore-5-(4-othony-5-methyl-1,3-thiazol-2-
y1) phenyl 1-methyl 6-(trifluoromethyl) -2, 4(1H, 3H)
pyrimidinedione;
-3-[4-chloro-2-fluoro-5-(4-mothoxy-5-methyl-1,3-thiozol-2-
yl) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3 [4-chlore-2-fluore-5-(4-cthoxy-5-mcthyl-1,3-thlazel-2-
yl)phenyl 1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinediene:
-3-[4-chloro-3-(4-benzyloxy-5-methyl-1,3-thiazol-2-
yl)phenyl-1-methyl-6-(trifluoremethyl)-2,4(1H,3H)-
pyrimidinedione,
- 3-[2,4-dichloro-5-(4-benzyloxy-5-methyl-1,3-thiazel-2-
v1) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione.
-3 [4-chloro-2-fluoro-5-(4-benzylony-5-methyl-1,3-thiazol-
2-yl)phenyl-1-mothyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-(2,4-dichloro-5-(5-(trifluoromethyl)-1,3,4-thiadiazel-
2-y1|oxy|pheny1)-6-(trifluoromethy1)-2,4(1H,3H)-
pyrimidinedioner
- 3-(4-chlore-2-fluore-5-([5-(trifluoremethyl)-1,3,4-
thiadiazel-2-yl]exy}phenyl)-6 (triflueromethyl)-2,4(1H,3H)-
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pyrimidinedione.
- 3-(2,4-dichlore-5-([5-(trifluoromethyl)-1,3,4-omadiazol-
2-yl|oxy|phonyl)-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedioner
- 3-(4-chloro-2-fluoro-5-(-5-(trifluoromethyl)-1,3,4-
exadiazel-2-yl]exy)phenyl)-6-(trifluoremethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4-chloro-3-({5-(trifluoremethyl)-1,3,4-thiadiazol-2-
vlloxvlphenyl) -1-methyl-6-(trifluoromethyl) -2,4(1H,3H)-
pyrimidinedione;
-3-(2.4 dichlore-5-(f5-(trifluoromethyl)-1,3,4-thiadiazel-
2-y1exy)phenyl)=1-methyl=6-(trifluoromethyl)=2,4(1H,3H)
pvrimidinedione.
- 3-(4-chloro-2-fluoro-5-([5-(trifluoromethyl)-1,3,4-
thiadiazel-2-yllexy)phenyl)-1-methyl-6-(txiflueromethyl)
2,1(1H,3H)-pyrimidinedioner
-3-(4-chloro-3-(5-methyl-1,3,4-thiadiazol-2-
yl) oxy]phenyl} l-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinediene;
-3-(2,4-dichlore-5-[(5-methyl-1,3,4-thiadiazel-2-
yl) oxy]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione:
- 3-(4-chlore-2-fluore-5-[(5-methyl-1,3,4-thiadiazel-2-
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y1) oxy]phonyl]-1-mcthyl-6-(trifluoromethyl)-2,4(1H,3H)-

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pyrimidinedione:
-3 (4-chloro-3-({5-(trifluoremethyl)-1,3,4-enadiazel-2-
yl]oxy)phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1K,3K)-
pyrimidinedione;
- 3-(2,4-dichlore-5-[[5-(trifluoremethyl)-1,3,4-exadiazel-
2-ylloxylphonyl) 1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4-chloro-3-fluoro-5-([5-(trifluoromothyl)-1,3,4-
omadiazol-2-yl]oxy)phenyl)-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H)-pyrimidinedione,
- 3-{4-chloro-3-{(5-methyl-1,3,4-enadlazol-2-
yl) oxylphonyl) -1-methyl-6-(trifluoromethyl) -2,4(1H,3H)-
pyrimidinedione+
- 3-(2,4-dichlore-5-[(5-methyl-1,3,4-oxadiazol-2-
yl) oxy)phonyl) -1-methyl-6-(trifluoromethyl) -2,4(1H,3H)-
pyrimidinedione,
- 3-(4-chlore-2 fluore-5-((5-methyl-1,3,4-emadiazel-2-
yl) oxy]phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinodione;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-6-oxo-2-thioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-{2-chloro-4-fluoro-5-{1,2,3,6-tetrahydro-3-
```

difluoromethyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-

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1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione; - 3-{4-chloro-3-[1-(ethoxymethyl)-1H-tetrazol-5-yl]phenyl}-

pyrimidinedione: - 3-{4-chloro-3-[2-(ethoxymethyl)-2H-tetrazol-5-yl]phenyl)-

y1] pheny1}-1-methy1-6-(trifluoromethy1)-2,4(1H,3H)-

pyrimidinedione; 3-{4-chloro-3-[1-(methoxymethyl)-1H-tetrazol-5-

 $3-\{4-chloro-3-[2-(methoxymethyl)-2H-tetrazol-5$ yl]phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-

methyl-2, 6-dioxo-4-(trifluoromethyl)pyrimidin-1yl]phenoxy}-3-methoxybut-2-enoate;

ethyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1vl]phenoxv}-3-methoxvbut-2-enoate;

ethyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-

methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1yl]phenoxy}-3-methoxypent-2-enoate;

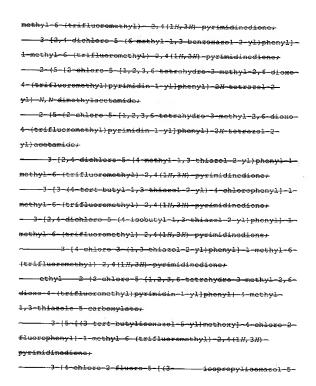
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1yl]phenoxy}-3-methoxypent-2-enoate; - methyl (2E)-4-(2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-

----3-[4-chloro-3-(4,5-dimethyl-1,3-thiazol-2-yl)phenyl]-1methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinediene; methy1 (2E) -4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-

yl]phenoxy}-3-methoxybut-2-enoate;

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1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[3-(2-ally1-2H-tetrazol-5-y1)-4-chloropheny1]-1-methyl-
 6-(trifluoromethyl) - 2,4(1H,3H)-pyrimidinedione;
- 3-[3-(1-allyl-1H-tetrazol-5-yl)-4-chlorophenyl]-1-methyl-
6-(trifluoromethyl) - 2,4(1H,3H)~pyrimidinedione;
         3-(4-chloro-2-fluoro-5-[(3-methyliooxazol-5-
y-1) methoxy | pheny1 | -1 - methy1 - 6 - (trifluoromethy1) - 2, 4 (1H, 3H) -
pyrimidinedione;
                  3 (2,4-dichloro-5-[(3-methylisoxazol-5-
yl)methoxylphenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione,
      --- 3-[4-chlore-3-(4-isopropoxy-5-methyl-1,3-thiazol-2-
yl)phenyl-1-methyl-6-(trifluoremethyl)-2,4(1H.3H)-
pyrimidinedione.
       3-[4-chlore-3-(4-hydrexy-5-methyl-1-3-thiazel-2-
yl)phonyl-1-methyl 6-(trifluoromethyl) 2,4(1H.3H)-
pyrimidinediene,
        3-{4-chloro-2-fluoro-5-[(5-methyl-1,2,4-oxadiazol-3-
yl)methoxy[phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
             3-(2,4-dichloro-5-[(5-methyl-1,2,4-oxadiazol-3-
y1) methoxy] phenyl) -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H) -
pyrimidinedione;
 3-[3-(1,3-benzothiazol-2-yl)-4-chlorophenyl]-1-methyl-6-
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(trifluoromethyl) - 2,4(1H,3H)-pyrimidinedione;
    - 3-[3-(1,3-benzoxazel-2-yl)-4-shlorophenyl]-1-methyl-6-
(trifluoromethyl) -2,4(1H,3H)-pyrimidinedione;
        3-{4-chloro-2-fluoro-5-[(3-methyl-1,2,4-oxadiazol-5-
yl)methoxy]phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-[1-chloro-3-(4-methyl-1,3-thiazol-2-yl)phenyl-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinediono;
                  3-[4-chloro-2-fluoro-5-(1,2,4-oxadiazol-3-
ylmethoxy) phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione:
     3-[3-(2-tert-butyl-2H-tetrazol-5-y1)-4-chlorophenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[5-(1,3-benzethiazel-2-yl)-4-chlore-2-fluerephenyl]-1-
methyl-6-(trifluoromethyl) - 2,4(1H,3H)-pyrimidinedioner
- 3-(4-chloro-3-{2-[(2-methoxyethoxy)methyl]-2H-tetrazol-5-
yl)phenyl) -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H) -
pvrimidinedione;
  3-(4-chloro-3-\{1-[(2-methoxyethoxy)methyl]-1H-tetrazol-5-
yl}phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-{5-(1,3-benzoxazol-2-yl)-4-chlore-2-fluorophenyl]-1-
methyl-6-(trifluoromethyl) 2,4(1H,3H)-pyrimidinedione;
       3-(5-(1,3-benzethiazel-2-yl)-2,4-dichlorophenyl)-1-
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y1) methoxy]pheny1}-1-methy1-6-(trifluoremethy1)-2,4(1H,3H)-
pyrimidinediene:
      3-[4-chloro-3-(2-isopropyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
         3-[3-(2-benzy1-2H-tetrazo1-5-y1)-4-chloropheny1]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
         3-[3-(1-benzyl-1H-tetrazol-5-yl)-4-chlorophenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
       3 (4-chloro-2-fluoro-5-((1-methyl-1H-tetrazol-5-
yl) exylphenyl) -1-methyl-6-(trifluoromethyl) -2,4(1H, 3H) -
pyrimidinedione;
        -- 3-(4-chlore-2-fluore-5-(4-methyl-2H-tetrazel-5-
yl) oxy]phenyl) - 1 methyl - 6 - (trifluoromethyl) - 2, 4 (1H, 3H) -
pvrimidinedione+
- methyl (2E)-4-{2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-dioxo-4(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
methoxybut-2-enoate;
    ethyl (2E)-4-{2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-dioxo-4(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
ethoxybut-2-enoate;
       3-[4-chloro-3-(1,2,4-oxadiazo1-3-ylmethoxy)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
```

```
-3-[4-chloro-3-(4,5,6,7-tetrahydro-1,3-benzothiazol-2-
yl) phenyl]-1-methyl-6-(trifluoremethyl)-2,4(14,34)-
pyrimidinedione+
---3-[4-chlore-3-(5,6-dihydro-1,4,2-dioxazin-3-yl)phenyl]-1-
mothyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione+
  exadiazin-2-yl)phenyl}-1-methyl-6-(trifluoremethyl)-
2,4(1H,3H)-pyrimidinedione;
-3-[1-chlore-3-(5,6-dihydre-1,1,2-diexagin-3-ylmethexyl-2-
fluorophenyl]-l-methyl-6 (trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione,
  3-(4-chloro-2-fluoro-5-(4-methyl-5-oxo-5-6-dihydro-4H-
1,3,4-oxadiazin-2-yl)methoxy]phenyl)-1-methyl-6-
(trifluoromothyl) -2, 4(1H, 3H) -pyrimidinodione;
 -----3-[4-chloro-3-(2-phonyl 2H-tetrazol-5-yl)phonyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
     3-[4-ohloro-3-(1-phenyl-1H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedioner
        3-(4-chloro-3-(1-(cyclopropylmethyl)-1H-tetrazol-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
        3-{4-chloro-3-[2-(cyclopropylmethyl)-2H-tetrazol-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
```

```
- 3-(4-chlore-3-[1-(2-exepropyl)-1H-tetrazel-5-yl]phenyl}-
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
  - 3-{4-chlore-3-{2-{2-exepropyl}-2#-tetrazel-5-yl]phenyl}-
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione.
- 3 [4-chlore-3-(4-evelopropyl-1,3 thiazel-2-yl)phenyl|-1
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione+
     3 (4-chloro-3-[4-(4-chlorophenyl)-1,3-thiazol-2-
vllphenyl]-1-methyl 6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione+
- ethyl - 2-{2-chloro-5-{1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo 1 (trifluoromothyl)pyrimidin-1-yl)phonyl)-1,3-
thiazolo-4-carboxylater
- 3-[3-(2-buty1-2H-tetrazol-5-y1)-4-chloropheny1]-1-methy1-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[4-chlore-2-fluore-5-(5,6-dihydre-1,4,2-diexamin-3-
vlmethoxv)-2-fluorophenyl]-1-methyl-6 (trifluoromethyl)-
2,4(1H,3H) pyrimidinedione,
- 3-(4-chloro-3-{2-[(4-chlorophenoxy)methyl]-2H-tetrazol-5-
yl)phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4-chloro-3-(1-[(4-chlorophenoxy)methyl]-1H-tetrazol-5-
yl}phenyl) -1-methyl-6-(trifluoromethyl) -2,4(1H,3H) -
pyrimidinedione;
  -3-[3-(4-tert-butyl-5-exe-4,5-dihydre-1,3,4-thiadiagel-2-
```

y1) 4-chlorephenyl]-1 methyl 6-(trifluoromethyl)2,1(1//,3//) pyrimidinedioner

- 3-{4-chloro-3-[2-(4-chlorobenzyl)-2H-tetrazol-5-yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)pyrimidinedione;
- 3-{4-chloro-3-[1-(4-chlorobenzyl)-1H-tetrazol-5-yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)pyrimidinedione;
- methyl 2-{2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1,3thiazole-4-carboxylate;
- methyl (2-(2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl)-1,3thiazol-4-vl)acetate.
- (canceled)
- 4. (withdrawn): A process for the preparation of compounds having general formula (I) according to claim 1, characterized in that it includes a cyclo-condensation reaction of an isocyanate or isothiocyanate having general formula (II) with a 3-aminocrotonate having general formula (III) according to reaction scheme 1

 Scheme 1:

$$X_2$$
 X_3
 X_4
 X_4
 X_2
 X_4
 X_5
 X_4
 X_5
 X_5
 X_5
 X_5
 X_6
 X_6
 X_6
 X_7
 X_8
 X_9
 X_9

wherein

- X_1 , X_2 , X_3 , X_4 , R and G have the meanings previously defined:
- R_{13} represents a C_1 - C_4 alkyl or C_1 - C_4 haloalkyl group or a phenyl group possibly substituted with C_1 - C_4 alkyl groups.
- 5. (withdrawn): The process according to claim 4, characterized in that the reaction is carried out in the presence of an inert organic solvent and in the presence of an organic base or preferably inorganic base, at a temperature ranging from -20°C to the boiling point of the reaction mixture.
- 6.(withdrawn): The process according to claim 4, characterized in that the isocyanates or isothiocyanates having general formula (II) are prepared starting from a

substituted aniline having general formula (IV) by reaction with a compound having general formula (V), such as phosgene, diphosgene, triphosgene or thiophosgene, according to reaction scheme 2

Scheme 2:

$$\begin{array}{c|c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
(V) & X_2
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_1 \\
X_2 & X_3
\end{array}$$

$$\begin{array}{c}
X_1 & X_2 & X_3
\end{array}$$

wherein

- X1, X2, X3 and G have the meanings defined above;
- L_3 and L_4 , the same or different, represent a chlorine atom or a CCl₃O- group.
- 7. (withdrawn): The process according to claim 6, characterized in that the reaction is carried out in the presence of an inert organic solvent, at a temperature ranging from 0°C to the boiling point of the mixture itself, possibly in the presence of a catalyst such as triethylamine, in an amount ranging from 0.001 and 100% by weight with respect to the aniline (IV), with a quantity of

reagent (V) varying from 1 to 3 moles per mole of aniline (IV).

8. (withdrawn): The process for the preparation of compounds having general formula (I) according to claim 1, wherein X_3 represents a $Q(CR_1R_2)_mZ^-$ group, a Q_1Z^- group, a $Y(OC)^-$ CR₈-CR₅-CR₅R₄Z⁻ group, compounds (Ia), characterized in that it comprises the reaction of a uracil having general formula (VI) with a compound having general formula (VII) according to reaction scheme 3

Scheme 3:

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wherein

- X1, X2, X4, G and Z have the meanings previously defined;
- R represents a $C_1\text{-}C_3$ alkyl group or a $C_1\text{-}C_3$ haloalkyl group;
- W represents a $Q(CR_1R_2)_n$ group, a Q_1 group, a X(OC)- CR_6 = CR_3 - CR_3R_6 group, wherein R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , Y, Q and Q_1 have the meanings defined above;
- L_2 represents a halogen atom, a R_LSO_2O- group, wherein R_L represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group or a phenyl group possibly substituted by C_1-C_4 alkyl groups, or it represents a $R_{L1}SO_2-$ group wherein R_{L1} represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group.
- 9. (withdrawn): The process according to claim 8, characterized in that the reaction between the compounds having general formula (VI) and the compounds having general formula (VII) is carried out in the presence of one or more inert organic solvent(s) and in the presence of a base, preferably an inorganic base, at a temperature ranging from -10°C to the boiling point of the reaction mixture.
- 10. (withdrawn): The process for the preparation of the compounds having general formula (I) according to claim 1, wherein G=0 and $R\neq H$, compounds (Ic), characterized in that it comprises the reaction of a uracil having general

formula (Ib) with an alkylating compound having general formula (VIII) according to reaction scheme 4

$$\begin{array}{c|c}
X_1 & & & & \\
X_2 & & & & \\
X_3 & & & & \\
\end{array}$$
(Ib) (Ic)

wherein

- X1, X2, X3 and X4 have the meanings defined above;
- R' represents a C1-C3 alkyl or C1-C3 haloalkyl group;
- L_1 represents a halogen atom, or a R_LSO_2O- group wherein \hat{R}_L represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group or a phenyl group possibly substituted by C_1-C_4 alkyl groups.
- 11. (previously presented): The process according to claim 10, characterized in that the reaction between the compounds having general formula (Ib) and the compound having general formula (VIII) is carried out in the presence of one or more inert organic solvents and in the presence of a base, preferably an inorganic base, at a temperature ranging from

- -10°C to the boiling point of the reaction mixture.
- 12. (withdrawn): The process according to claim 8, characterized in that the reaction is carried out in a biphasic system using water as solvent and an organic solvent immiscible with water, in the presence of phase transfer catalysts.
- 13. (withdrawn): The process for the preparation of compounds having general formula (I) according to claim 1, wherein G=0, compounds (Id), characterized in that it comprises a first reaction between a substituted aniline having formula (IV) and a chloroformiate or a carbonate having formula (IX) to give a carbamate having formula (X) and a second reaction wherein the carbamate is converted into the compounds having general formula (Id) by cyclocondensation with a 3-aminocrotonate having general formula (III), according to reaction scheme 5:

Scheme 5:

wherein

- X1, X2, X3, X4 and R have the meanings defined above;
- L₅ represents a halogen atom or a OR₁₄ group;
- R_{13} and R_{14} represent a C_1 - C_4 alkyl or C_1 - C_6 haloalkyl group or a phenyl group possibly substituted by C_1 - C_4 alkyl groups.
- 14. (withdrawn): The process according to claim 13, characterized in that the first reaction is carried out in the presence of an inert organic solvent, at a temperature ranging from -10°C to the boiling point of the mixture

itself, in the presence of an organic or inorganic base, in a quantity varying from 1 to 1.5 moles per mole of aniline (IV), with a quantity of compound having formula (IX) varying from 1 to 1.5 moles per mole of aniline (IV).

15. (withdrawn): The process according to claim 13, characterized in that the cyclo-condensation reaction of the carbamate having general formula (X) with the 3-aminocrotonate having general formula (III) is carried out in the presence of an inert organic solvent and in the presence of an organic or prefexably inorganic base, at a temperature ranging from -20°C to the boiling point of the reaction mixture.

16. (withdrawn): The process according to claim 10, characterized in that the compounds having general formula (Ib) are prepared starting from an aniline having general formula (IV) by reaction with a β -ketoester having general formula (XII), to give an anilide having general formula (XIII), then converted into the intermediate of general formula (XIV) by amination with ammonia or ammonium salts, said intermediate being converted into the compounds of general formula (Ib) by cyclization with a compound of general formula (XV), such as phosgene, or diphosgene according to the reaction scheme 6

Scheme 6:

wherein:

- X1, X2, X3 and X4 have the meanings defined above;
- R_{13} represents a C_1 - C_4 alkyl or haloalkyl group or a phenyl group possibly substituted by C_1 - C_4 alkyl groups;
- L_6 and L_7 , having the same or different meaning, represent
- a chlorine atom, a CCl_2O- group, a C_1-C_4 alkoxy group, a phenoxy group, an imidazol-1-yl group or a 1,2,4-triazol-1-yl group.
- 17. (withdrawn): The process according to claim 16,

characterised in that the reaction between the compounds having general formula (IV) and the compounds having general formula (XII) is carried out in the presence of one or more inert organic solvents, at a temperature ranging from -10°C to the boiling temperature of the reaction mixture, using an amount of compound (XII) ranging from 1 to 3 moles per mole of aniline (IV).

- 18. (withdrawn): (currently amended) The process according to claim 17, characterised in that the reaction is carried out while distilling off compound $R_{13}\mathrm{OH}$ formed during the reaction, alone or in mixture with the solvent used.
- 19. (withdrawn): (currently amended) The process according to claim 16, characterised in that the transformation of compounds having general formula (XIII) into compounds having general formula (XIV) is carried out in the presence of one or more inert organic solvents, at a temperature ranging from -10°C to the boiling temperature of the reaction mixture, using ammonia or an ammonium salt, in an amount ranging from 1 to 20 moles per mole of compound (XIII).
- 20. (withdrawn): The process according to claim 16, characterised in that the reaction between the compounds having general formula (XIV) and the compounds having general formula (XV) is carried out in the presence of one-

or more inert organic solvents, at a temperature ranging from $-10\,^{\circ}\text{C}$ to the boiling temperature of the reaction mixture, using an amount of compound (XV) ranging from 1 to 5 moles per mole of compound (XIV) in the presence of a suitable organic or inorganic base, in an amount ranging from 1 to 5 moles per mole of compound (XIV).

- 21. (withdrawn): Use of uracils having general formula
 (I) according to claims 1, as herbicides.
 - 22.(withdrawn): Use according to claim 21 for the preemergence and/or post-emergence control of monocotyledonous or dicotyledonous weeds.
 - 23. (withdrawn): Method for the control of weeds in cultivated areas by the application of the compounds having general formula (I) according to claims 1.
 - 24. (withdrawn): (The method according to claim 23, characterized in that the amount of compound having formula (I) to be applied varies between dosages of compounds ranging from 1g to 1000g per hectare.
 - 25.(currently amended): The herbisidal composition[[s]] containing, as active principle, one or more compounds

having general formula (I) according to claim 1, possibly also as a blend of isomers.

- 26. (currently amended): The herbicidal composition[[s]] according to claim 25, comprising other active principles which are compatible with the compounds having general formula (I), such as and are selected from the group consisting of other herbicides, fungicides, insecticides, acaricides, and fertilizers, etc.
- 27. (currently amended): The herbicidal composition[[s]] according to claim [[25]] 26, characterized in that the further other herbicides are selected from: acetochlor, acifluorfen, aclonifen, AKH-7088, alachlor, alloxydim, ametryn, amicarbazone, amidosulfuron, amitrole, anilofos, asulam, atrazine, azafenidin, azimsulfuron, aziprotryne, BAY MKH 6561, beflubutamid, benazolin, benfuresate, bensulfuron, benfluralin, bensulide. bentazone, benzfendizone, benzobicyclon, benzofenap, benzthiazuron, bifenox, bilanafos, bispyribac-sodium, bromacil, bromobutide, bromofenoxim, bromoxynil, butachlor, butafenacil, butamifos, butenachlor, butralin, butroxydim, butylate, cafenstrole, carbetamide, carfentrazone-ethyl, chlomethoxyfen, chloramben, chlorbromuron, chlorbufam,

chlorflurenol, chloridazon, chlorimuron, chlornitrofen, chlorotoluron, chloroxuron, chlorpropham, chlorsulfuron, chlorthal, chlorthiamid, cinidon ethyl, cinmethylin, cinosulfuron, clethodim, clodinafop, clomazone, clomeprop, clopyralid, cloransulam-methyl, cumyluron (JC-940), cyanazine, cycloate, cyclosulfamuron, cycloxydim, cyhalofop-butyl, 2,4-D, 2,4-DB, daimuron, dalapon, desmedipham, desmetryn, dicamba, dichlobenil, dichlorprop, dichlorprop-P, diclofop, diclosulam, diethatyl, difenoxuron, difenzoquat, diflufenican, diflufenzopyr, dimefuron, dimepiperate, dimethachlor, dimethametryn, dimethenamid, dinitramine, dinoseb, dinoseb acetate, dinoterb, diphenamid, dipropetryn, diquat, dithiopyr, 1diuron, eglinazine, endothal, EPTC, esprocarb, ethalfluralin, ethametsulfuron-methyl, ethidimuron, ethiozin (SMY 1500), ethofumesate, ethoxyfen-ethyl (HC-252), ethoxysulfuron, etobenzanid (HW 52), fenoxaprop, fenoxaprop-P, fentrazamide, fenuron, flamprop, flamprop-M, flazasulfuron, florasulam, fluazifop, fluazifop-P, fluazolate (JV 485), flucarbazone-sodium, fluchloralin, flufenacet, flufenpyr ethyl, flumetsulam, flumicloracpentyl, flumioxazin, flumipropin, fluometuron, fluoroglycofen, fluoronitrofen, flupoxam, flupropanate, flupyrsulfuron, flurenol, fluridone, flurochloridone,

fluroxypyr, flurtamone, fluthiacet-methyl, fomesafen, foramsulfuron, fosamine, furyloxyfen, glufosinate, glyphosate, halosulfuron-methyl, haloxyfop, haloxyfop-Pmethyl, hexazinone, imazamethabenz, imazamox, imazapic, imazapyr, imazaguin, imazethapyr, imazosulfuron, indanofan, iodosulfuron, ioxynil, isopropalin, isoproturon, isouron, isoxaben, isoxachlortole, isoxaflutole, isoxapyrifop, KPP-421, lactofen, lenacil, linuron, LS830556, MCPA, MCPAthioethyl, MCPB, mecoprop, mecoprop-P, mefenacet, mesosulfuron, mesotrione, metamitron, metazachlor, methabenzthiazuron, methazole, methoprotryne, methyldymron, metobenzuron, metobromuron, metolachlor, S-metolachlor, metosulam, metoxuron, metribuzin, metsulfuron, molinate, monalide, monolinuron, naproanilide, napropamide, naptalam, NC-330, neburon, nicosulfuron, nipyraclofen, norflurazon, orbencarb, oryzalin, oxadiargyl, oxadiazon, oxasulfuron, oxaziclomefone, oxyfluorfen, paraquat, pebulate, pendimethalin, penoxsulam, pentanochlor, pentoxazone, pethoxamid, phenmedipham, picloram, picolinafen, piperophos, pretilachlor, primisulfuron, prodiamine, profluazol, proglinazine, prometon, prometryne, propachlor, propanyl, propaquizafop, propazine, propham, propisochlor, propyzamide, prosulfocarb, prosulfuron, pyraclonil, pyraflufen-ethyl, pyrazogyl (HSA-961), pyrazolynate,

pyrazosulfuron, pyrazoxyfen, pyribenzoxim, pyributicarb, pyridafol, pyridate, pyriftalid, pyriminobac-methyl, pyrithiobac-sodium, quinclorac, quinmerac, quizalofop, quizalofop-P, rimsulfuron, sethoxydim, siduron, simazine, simetryn, sulcotrione, sulfentrazone, sulfometuron-methyl, sulfosulfuron, 2,3,6-TBA, TCA-sodium, tebutam, tebuthiuron, tepraloxydim, terbacil, terbumeton, terbuthyl-azine, terbutryn, thenylchlor, thiazafluron, thiazopvr, thifensulfuron-methyl, thidiazimin. thiobencarb, tiocarbazil, tioclorim, tralkoxydim, tri-allate, triasulfuron, triaziflam, tribenuron. triclopyr, trietazine, trifloxysulfuron, trifluralin, triflusulfuronmethyl, tritosulfuron, UBI-C4874, vernolate.

- 28. (currently amended): The composition[[s]] according to claim 25, characterized in that the concentration of the active substance ranges from 1 to 90%.
- 29.(new): A uracil compound as defined in claim 1 wherein Q is 1,2,4-oxadiazoly1.
- 30.(new): A uracil compound as defined in claim 1 wherein Q is 5-methyl-1,2,4-oxadiazolyl.